



General Assembly

Substitute Bill No. 7124

January Session, 2017

* _____HB07124INS_____031017_____*

AN ACT CONCERNING MAXIMUM ALLOWABLE COST LISTS AND DISCLOSURES BY PHARMACY BENEFITS MANAGERS, LIMITING COST-SHARING FOR PRESCRIPTION DRUGS AND SHIELDING PHARMACISTS AND PHARMACIES FROM CERTAIN PENALTIES.

Be it enacted by the Senate and House of Representatives in General Assembly convened:

1 Section 1. (NEW) (*Effective October 1, 2017*) (a) As used in this
2 section, (1) "maximum allowable cost" means the maximum amount a
3 pharmacy benefits manager will reimburse a pharmacy for a
4 prescription drug, and (2) "maximum allowable cost list" means a list
5 of prescription drugs for which a maximum allowable cost has been
6 established by a pharmacy benefits manager.

7 (b) (1) Each pharmacy benefits manager shall, prior to placing a
8 prescription drug on a maximum allowable cost list, ensure that such
9 drug (A) (i) has been designated as therapeutically equivalent to other
10 pharmaceutically equivalent products with an "A" code or "B" code in
11 the most recent edition or supplement of the federal Food and Drug
12 Administration's Approved Drug Products with Therapeutic
13 Equivalence Evaluations, or (ii) has an "NR" rating, "NA" rating or
14 similar rating by a nationally recognized pricing reference, and (B) (i)
15 is available for purchase by pharmacies in this state from national or
16 regional wholesalers, and (ii) is not obsolete or temporarily
17 unavailable. As used in this subparagraph, a drug is obsolete even if it

18 is listed in national drug pricing compendia, if it is no longer actively
19 marketed by the manufacturer or labeler.

20 (2) Each pharmacy benefits manager shall remove a prescription
21 drug from a maximum allowable cost list not later than three business
22 days after (A) the prescription drug no longer meets the requirements
23 in subdivision (1) of this subsection, or (B) the pharmacy benefits
24 manager becomes aware that such drug no longer meets the
25 requirements in subdivision (1) of this subsection.

26 (c) Each contract entered into, renewed or amended on or after
27 October 1, 2017, between a pharmacy benefits manager and a
28 pharmacy or a pharmacy's contracting representative or agent shall
29 disclose the sources used by the pharmacy benefits manager to
30 determine the maximum allowable costs for prescription drugs on
31 each maximum allowable cost list for such pharmacy.

32 (d) Each pharmacy benefits manager shall:

33 (1) Provide an updated maximum allowable cost list to a plan
34 sponsor whenever there is a change to any such list under the plan;

35 (2) Update each maximum allowable cost list at least every seven
36 calendar days and promptly notify and make available to each in-
37 network pharmacy any such updated list applicable to such pharmacy;
38 and

39 (3) Establish an appeals process for a pharmacy to contest the
40 maximum allowable cost of a prescription drug in accordance with the
41 provisions of subsection (e) of this section. Each pharmacy benefits
42 manager shall provide to each in-network pharmacy information
43 concerning the appeals process.

44 (e) (1) A pharmacy may contest the maximum allowable cost of a
45 prescription drug based on one or both of the following grounds:

46 (A) The prescription drug does not meet the requirements in

47 subdivision (1) of subsection (b) of this section; or

48 (B) The maximum allowable cost established by the pharmacy
49 benefits manager for the prescription drug is below the cost at which
50 such drug is available for purchase from national or regional
51 wholesalers.

52 (2) A pharmacy contesting the maximum allowable cost of a
53 prescription drug shall file an appeal with the pharmacy benefits
54 manager not later than sixty calendar days after filing its submission
55 for the initial claim for reimbursement for such drug. The pharmacy
56 benefits manager shall investigate and issue a determination of such
57 appeal not later than seven calendar days after such manager receives
58 such appeal.

59 (3) If the pharmacy benefits manager determines the appeal is
60 denied, the manager shall provide to the pharmacy the reason for the
61 denial and the national drug code of a therapeutically equivalent
62 prescription drug that is available for purchase by pharmacies in this
63 state from national or regional wholesalers at a price that is equal to or
64 less than the maximum allowable cost for the prescription drug that is
65 the subject of the appeal.

66 (4) If the pharmacy benefits manager determines the appeal is valid,
67 such manager shall (A) adjust the maximum allowable cost for such
68 prescription drug, and (B) adjust such maximum allowable cost for the
69 appealing pharmacy not later than five business days after making
70 such determination.

71 Sec. 2. Section 38a-510 of the general statutes is repealed and the
72 following is substituted in lieu thereof (*Effective January 1, 2018*):

73 (a) No insurance company, hospital service corporation, medical
74 service corporation, health care center or other entity delivering,
75 issuing for delivery, renewing, amending or continuing an individual
76 health insurance policy or contract that provides coverage for
77 prescription drugs may:

78 (1) Require any person covered under such policy or contract to
79 obtain prescription drugs from a mail order pharmacy as a condition
80 of obtaining benefits for such drugs; [or]

81 (2) Impose a coinsurance, copayment, deductible or other out-of-
82 pocket expense that exceeds the claim cost of a covered prescription
83 drug, except that a high deductible health plan, as that term is used in
84 subsection (f) of section 38a-493, shall not be subject to the deductible
85 provision set forth in this subdivision until after the minimum annual
86 deductible for such plan has been met; or

87 [(2)] (3) Require, if such insurance company, hospital service
88 corporation, medical service corporation, health care center or other
89 entity uses step therapy for such drugs, the use of step therapy for any
90 prescribed drug for longer than sixty days. At the expiration of such
91 time period, an insured's treating health care provider may deem such
92 step therapy drug regimen clinically ineffective for the insured, at
93 which time the insurance company, hospital service corporation,
94 medical service corporation, health care center or other entity shall
95 authorize dispensation of and coverage for the drug prescribed by the
96 insured's treating health care provider, provided such drug is a
97 covered drug under such policy or contract. If such provider does not
98 deem such step therapy drug regimen clinically ineffective or has not
99 requested an override pursuant to subdivision (1) of subsection (b) of
100 this section, such drug regimen may be continued. For purposes of this
101 section, "step therapy" means a protocol or program that establishes
102 the specific sequence in which prescription drugs for a specified
103 medical condition are to be prescribed.

104 (b) (1) Notwithstanding the sixty-day period set forth in subdivision
105 [(2)] (3) of subsection (a) of this section, each insurance company,
106 hospital service corporation, medical service corporation, health care
107 center or other entity that uses step therapy for such prescription
108 drugs shall establish and disclose to its health care providers a process
109 by which an insured's treating health care provider may request at any
110 time an override of the use of any step therapy drug regimen. Any

111 such override process shall be convenient to use by health care
112 providers and an override request shall be expeditiously granted when
113 an insured's treating health care provider demonstrates that the drug
114 regimen required under step therapy (A) has been ineffective in the
115 past for treatment of the insured's medical condition, (B) is expected to
116 be ineffective based on the known relevant physical or mental
117 characteristics of the insured and the known characteristics of the drug
118 regimen, (C) will cause or will likely cause an adverse reaction by or
119 physical harm to the insured, or (D) is not in the best interest of the
120 insured, based on medical necessity.

121 (2) Upon the granting of an override request, the insurance
122 company, hospital service corporation, medical service corporation,
123 health care center or other entity shall authorize dispensation of and
124 coverage for the drug prescribed by the insured's treating health care
125 provider, provided such drug is a covered drug under such policy or
126 contract.

127 (c) Nothing in this section shall (1) preclude an insured or an
128 insured's treating health care provider from requesting a review under
129 sections 38a-591c to 38a-591g, inclusive, or (2) affect the provisions of
130 section 38a-492i.

131 (d) No individual health insurance carrier may terminate the
132 services of, require additional documentation from, require additional
133 utilization review, reduce payments or otherwise penalize or provide
134 financial disincentives to any pharmacy or pharmacist on the basis that
135 the pharmacy or pharmacist disclosed to an insured information
136 concerning (1) the cost or efficacy of a prescription drug, or (2) any
137 drug that is therapeutically equivalent to a prescription drug.

138 Sec. 3. Section 38a-544 of the general statutes is repealed and the
139 following is substituted in lieu thereof (*Effective January 1, 2018*):

140 (a) No insurance company, hospital service corporation, medical
141 service corporation, health care center or other entity delivering,

142 issuing for delivery, renewing, amending or continuing a group health
143 insurance policy or contract that provides coverage for prescription
144 drugs may:

145 (1) Require any person covered under such policy or contract to
146 obtain prescription drugs from a mail order pharmacy as a condition
147 of obtaining benefits for such drugs; [or]

148 (2) Impose a coinsurance, copayment, deductible or other out-of-
149 pocket expense that exceeds the claim cost of a covered prescription
150 drug, except that a high deductible health plan, as that term is used in
151 subsection (f) of section 38a-493, shall not be subject to the deductible
152 provision set forth in this subdivision until after the minimum annual
153 deductible for such plan has been met; or

154 [(2)] (3) Require, if such insurance company, hospital service
155 corporation, medical service corporation, health care center or other
156 entity uses step therapy for such drugs, the use of step therapy for any
157 prescribed drug for longer than sixty days. At the expiration of such
158 time period, an insured's treating health care provider may deem such
159 step therapy drug regimen clinically ineffective for the insured, at
160 which time the insurance company, hospital service corporation,
161 medical service corporation, health care center or other entity shall
162 authorize dispensation of and coverage for the drug prescribed by the
163 insured's treating health care provider, provided such drug is a
164 covered drug under such policy or contract. If such provider does not
165 deem such step therapy drug regimen clinically ineffective or has not
166 requested an override pursuant to subdivision (1) of subsection (b) of
167 this section, such drug regimen may be continued. For purposes of this
168 section, "step therapy" means a protocol or program that establishes
169 the specific sequence in which prescription drugs for a specified
170 medical condition are to be prescribed.

171 (b) (1) Notwithstanding the sixty-day period set forth in subdivision
172 [(2)] (3) of subsection (a) of this section, each insurance company,
173 hospital service corporation, medical service corporation, health care

center or other entity that uses step therapy for such prescription drugs shall establish and disclose to its health care providers a process by which an insured's treating health care provider may request at any time an override of the use of any step therapy drug regimen. Any such override process shall be convenient to use by health care providers and an override request shall be expeditiously granted when an insured's treating health care provider demonstrates that the drug regimen required under step therapy (A) has been ineffective in the past for treatment of the insured's medical condition, (B) is expected to be ineffective based on the known relevant physical or mental characteristics of the insured and the known characteristics of the drug regimen, (C) will cause or will likely cause an adverse reaction by or physical harm to the insured, or (D) is not in the best interest of the insured, based on medical necessity.

(2) Upon the granting of an override request, the insurance company, hospital service corporation, medical service corporation, health care center or other entity shall authorize dispensation of and coverage for the drug prescribed by the insured's treating health care provider, provided such drug is a covered drug under such policy or contract.

(c) Nothing in this section shall (1) preclude an insured or an insured's treating health care provider from requesting a review under sections 38a-591c to 38a-591g, inclusive, or (2) affect the provisions of section 38a-518i.

(d) No group health insurance carrier may terminate the services of, require additional documentation from, require additional utilization review, reduce payments or otherwise penalize or provide financial disincentives to any pharmacy or pharmacist on the basis that the pharmacy or pharmacist disclosed to an insured information concerning (1) the cost or efficacy of a prescription drug, or (2) any drug that is therapeutically equivalent to a prescription drug.

Sec. 4. Section 38a-479aaa of the general statutes is repealed and the

206 following is substituted in lieu thereof (*Effective October 1, 2017*):

207 As used in this section and sections 38a-479bbb to 38a-479iii,
208 inclusive, and section 1 of this act:

209 (1) "Commissioner" means the Insurance Commissioner;

210 (2) "Department" means the Insurance Department;

211 (3) "Drug" means drug, as defined in section 21a-92;

212 (4) "Person" means person, as defined in section 38a-1;

213 (5) "Pharmacist services" includes (A) drug therapy and other
214 patient care services provided by a licensed pharmacist intended to
215 achieve outcomes related to the cure or prevention of a disease,
216 elimination or reduction of a patient's symptoms, and (B) education or
217 intervention by a licensed pharmacist intended to arrest or slow a
218 disease process;

219 (6) "Pharmacist" means an individual licensed to practice pharmacy
220 under section 20-590, 20-591, 20-592 or 20-593, and who is thereby
221 recognized as a health care provider by the state of Connecticut;

222 (7) "Pharmacy" means a place of business where drugs may be sold
223 at retail and for which a pharmacy license has been issued to an
224 applicant pursuant to section 20-594; and

225 (8) "Pharmacy benefits manager" or "manager" means any person
226 that administers the prescription drug, prescription device, pharmacist
227 services or prescription drug and device and pharmacist services
228 portion of a health benefit plan on behalf of plan sponsors such as self-
229 insured employers, insurance companies, labor unions and health care
230 centers.

231 Sec. 5. Section 38a-479hhh of the general statutes is repealed and the
232 following is substituted in lieu thereof (*Effective October 1, 2017*):

233 (a) The commissioner may conduct investigations and hold hearings
 234 on any matter under the provisions of sections 38a-479aaa to 38a-479iii,
 235 inclusive, as amended by this act, or section 1 of this act. The
 236 commissioner may issue subpoenas, administer oaths, compel
 237 testimony and order the production of books, records and documents.
 238 If any person refuses to appear, to testify or to produce any book,
 239 record, paper or document when so ordered, upon application of the
 240 commissioner, a judge of the Superior Court may make such order as
 241 may be appropriate to aid in the enforcement of this section.

242 (b) Any person aggrieved by an order or decision of the
 243 commissioner under sections 38a-479aaa to 38a-479iii, inclusive, as
 244 amended by this act, or section 1 of this act may appeal therefrom in
 245 accordance with the provisions of section 4-183.

This act shall take effect as follows and shall amend the following sections:		
Section 1	<i>October 1, 2017</i>	New section
Sec. 2	<i>January 1, 2018</i>	38a-510
Sec. 3	<i>January 1, 2018</i>	38a-544
Sec. 4	<i>October 1, 2017</i>	38a-479aaa
Sec. 5	<i>October 1, 2017</i>	38a-479hhh

INS *Joint Favorable Subst.*